Attorney Docket No. 9310.28CT

In re: Goudsmit et al. Serial No.: 09/760,085 Filed: January 12, 2001

IN THE CLAIMS

Please amend the claims as follows.

1-15. (Canceled)

16. (Currently amended) A method for separating single stranded nucleic acid from double stranded nucleic acid, comprising the steps of:

contacting a mixture comprising both single stranded nucleic acid and double stranded nucleic acid with a first liquid comprising a chaotropic agent and a nucleic acid binding solid phase in the absence of material containing alcohol groups, wherein the first liquid has a composition such that the double stranded nucleic acid binds to the solid phase;

separating the solid phase from a supernatant containing the single stranded nucleic acid; and

contacting the supernatant with a second liquid comprising a second nucleic acid binding solid phase, in the presence of a chaotropic agent, and in the absence of material containing alcohol-groups wherein the second liquid has a composition such that the resulting mixture of supernatant and second liquid allows for binding of the single stranded nucleic acid to the second solid phase.

- 17. (Previously presented) The method according to claim 16, wherein the first liquid comprises a chaotropic agent in concentration between about 1-10M, and a chelating agent, and has a pH between about 2 and 10.
- 18. (Previously presented) The method according to claim 17, wherein the chelating agent is EDTA, which is present in a concentration between about 10 mM and 1 M.
- 19. (Previously presented) The method according to claim 18, wherein the first liquid comprises at least about 100 mM EDTA and guanidinium salt as a chaotropic agent.

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- 20. (Previously presented) The method according to claim 16, wherein the chaotropic agent is guanidinium thiocyante.
- 21. (Previously presented) A method according to claim 20, whereby the first liquid has the constitution of a buffer prepared by dissolving about 120g guianidinium thiocyanate in about 100ml 0.2M EDTA (pH=8).

22-27. (Canceled)

- 28. (Previously presented) The method according to claim 16, wherein the solid phase is silicum based.
- 29. (Previously presented) The method according to claim 28, wherein the solid phase is silica.
- 30. (Previously presented) The method according to claim 29, wherein the silica is in the form of particles having a size between about 0.05 and about 500 micrometers.
- 31. (Previously presented) The method according to claim 16, wherein the solid phase is separated from the supernatant by centrifugation.

32-37. (Canceled)

38. (Currently amended) A method for separating single stranded nucleic acid from double stranded nucleic acid, comprising the steps of:

contacting a mixture comprising both single stranded and double stranded nucleic acid with a first liquid comprising a chaotropic agent and a nucleic acid binding solid phase in the absence of material containing alcohol groups, wherein the first liquid has a composition such that the double stranded nucleic acid binds to the solid phase;

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separating the solid phase from a supernatant containing the single stranded nucleic acid; and

contacting the supernatant with a second liquid comprising a second nucleic acid binding solid phase, in the presence of a chaotropic agent, a chelating agent and divalent positive ionsand in the absence of material containing alcohol groups, wherein the second liquid has a composition such that the resulting mixture of supernatant and second liquid allows for binding of the single stranded nucleic acid to the second solid phase.

- 39. (Previously presented) The method according to Claim 38, wherein the concentration of the divalent positive ions is the same as the concentration of the chelating agent.
- 40. (Previously presented) The method according to Claim 38, wherein the chelating agent is EDTA and the ions are Mg²⁺ ions.
- 41. (Previously presented) The method according to Claim 38, wherein the chaotropic agent is a guanidinium salt.
- 42. (Previously presented) The method according to Claim 41, wherein the guanidinium salt is guanidinium isothiocyanate.
- 43. (Previously presented) The method according to Claim 42, wherein the second liquid has the constitution of a buffer prepared by dissolving about 120g guanidinium isothiocyanate in about 100ml 0.35M TRIS HCl (pH 6.4) and adding about 22ml 0.2 M EDTA (pH 8.0) and about 9.1g Triton X-100TM (polyethoxylated p-isooctyl-phenol), homogenizing the solution and adding MgCl₂ to a final concentration of about 0.25M.
- 44. (Currently amended) A method for separating single stranded nucleic acid from double stranded nucleic acid, comprising the steps of:

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contacting a mixture comprising both single stranded nucleic acid and double stranded nucleic acid with a first liquid comprising a chaotropic agent and a nucleic acid binding solid phase in the absence of material containing alcohol groups, wherein the first liquid has a composition such that the double stranded nucleic acid binds to the solid phase;

separating the solid phase from a supernatant containing the single stranded nucleic acid; and

contacting the supernatant with a second liquid comprising a second nucleic acid binding solid phase, in the presence of a chaotropic agent and divalent positive ions and in the absence of material containing alcohol groups, wherein the second liquid has a composition such that the resulting mixture of supernatant and second liquid allows for binding of the single stranded nucleic acid to the second solid phase.